ESIKA TOTAL SEC ULTRA FRESH- aluminum sesquichlorohydrate liquid
Ventura Corporation, Ltd.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ésika total sec ULTRA FRESH ALCOHOL-FREE MEN'S ROLL-ON DEODORANT AND ANTIPERSPIRANT ALL DAY EFFECTIVE PROTECTION

Drug Facts

Active Ingredient

Aluminum sesquichlorohydrate 16.2 %

Purpose

Antiperspirant

Uses

- Reduces underarm perspiration
- All day extra effective protection

Warnings

For external use only

Do not use on broken skin

Ask a doctor before use if you have Kidney desease

Stop use and ask a doctor if rash or irritation occurs

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply to underarms only.

Inactive ingredients

AQUA (WATER), STEARETH-2, STEARETH-21, PPG-15 STEARYL ETHER, CYCLOPENTASILOXANE, CYCLOHEXASILOXANE, PARFUM (FRAGRANCE), DICAPRYLYL CARBONATE, BISABOLOL, TRICLOSAN, BENZALKONIUM CHLORIDE, METHYLPARABEN, BHT, PROPYLENE GLYCOL, MENTHYL PCA, TETRASODIUM EDTA, MENTHOL, DIPROPYLENE GLYCOL, BAMBUSA ARUNDINACEA LEAF EXTRACT

PR: Dist. By Ventura Corporation, Ltd. San Juan, Puerto Rico 00926.

PRINCIPAL DISPLAY PANEL - 50 ml Bottle Label

ésika
total
sec

ULTRA FRESH

Alcohol-free
Men's Roll-on
Deodorant and
Antiperspirant
All day extra
effective protection

36

50 ml e (1.7 fl.oz.)

ESIKA TOTAL SEC ULTRA FRESH

aluminum sesquichlorohydrate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:13537-156
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM SESQUICHLOROHYDRATE (UNII: UCN889409V) (ALUMINUM SESQUICHLOROHYDRATE - UNII:UCN889409V)	ALUMINUM SESQUICHLOROHYDRATE	0.0162 g in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-21 (UNII: 53J3F32P58)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
LEVOMENOL (UNII: 24WE03BX2T)	
TRICLOSAN (UNII: 4NM5039Y5X)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

METHYLPARABEN (UNII: A2I8C7HI9T)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
MENTHYL PYRROLIDONE CARBOXYLATE, (-),DL- (UNII: 8P18J856U2)	
EDETATE SODIUM (UNII: MP1J8420LU)	
MENTHOL (UNII: L7T10EIP3A)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
BAMBUSA ARUNDINACEA LEAF (UNII: HW86D1FGSS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13537-156-01	50 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part350	01/24/2012	

Labeler - Ventura Corporation, Ltd. (602751344)

Establishment

Name	Address	ID/FEI	Business Operations
Bel Star S.A. (Colombia)		880160197	MANUFACTURE

Revised: 1/2012 Ventura Corporation, Ltd.